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NEWS 11 Apr 14 MEDLINE Reload
NEWS 12 Apr 17 Polymer searching in REGISTRY enhanced
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NEWS 19 May 19 Simultaneous left and right truncation added to WSCA
NEWS 20 May 19 RAPRA enhanced with new search field, simultaneous left and right truncation
NEWS 21 Jun 06 Simultaneous left and right truncation added to CBNB
NEWS 22 Jun 06 PASCAL enhanced with additional data
NEWS 23 Jun 20 2003 edition of the FSTA Thesaurus is now available
NEWS 24 Jun 25 HSDB has been reloaded
NEWS 25 Jul 16 Data from 1960-1976 added to RDISCLOSURE
NEWS 26 Jul 21 Identification of STN records implemented
NEWS 27 Jul 21 Polymer class term count added to REGISTRY
NEWS 28 Jul 22 INPADOC: Basic index (/BI) enhanced; Simultaneous Left and Right Truncation available
NEWS 29 AUG 05 New pricing for EUROPATFULL and PCTFULL effective August 1, 2003
NEWS 30 AUG 13 Field Availability (/FA) field enhanced in BEILSTEIN
NEWS 31 AUG 15 PATDPAFULL: one FREE connect hour, per account, in September 2003
NEWS 32 AUG 15 PCTGEN: one FREE connect hour, per account, in September 2003
NEWS 33 AUG 15 RDISCLOSURE: one FREE connect hour, per account, in September 2003
NEWS 34 AUG 15 TEMA: one FREE connect hour, per account, in September 2003
NEWS 35 AUG 18 Data available for download as a PDF in RDISCLOSURE
NEWS 36 AUG 18 Simultaneous left and right truncation added to PASCAL
NEWS 37 AUG 18 FROSTI and KOSMET enhanced with Simultaneous Left and Right Truncation
NEWS 38 AUG 18 Simultaneous left and right truncation added to ANABSTR

NEWS EXPRESS April 4 CURRENT WINDOWS VERSION IS V6.01a, CURRENT
MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
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=> s imatinib(w)mesylate?
L1 810 IMATINIB (W) MESYLAATE?

=> s 11 and leukemia?
L2 655 L1 AND LEUKEMIA?

=> s decitabine?
L3 208 DECITABINE?

=> s 12 and 13
L4 7 L2 AND L3

=> d 14 abs ibib 1-7

L4 ANSWER 1 OF 7 MEDLINE on STN

AB Chronic myeloid leukemia (CML) typically runs a biphasic or triphasic course, with diagnoses usually made in the chronic phase (CP). Without effective treatment, patients eventually progress to a blastic phase (BP), frequently through an intermediate or accelerated phase (AP). Because the definition of AP varies among studies, comparisons of outcome and prognosis are difficult. The management of patients in these advanced phases of the disease has been much less satisfactory than that of patients in CP. Treatment with interferon-alfa (IFNalpha)-based therapy is ineffective for most patients in AP and for all of those in BP.

Imatinib mesylate has demonstrated significant activity in AP and BP disease, although the results are inferior compared to treatment in CP. In AP, 82% of patients achieve a hematologic response, with 24% achieving a major cytogenetic remission (MCR). Early MCR (within 3 months of diagnosis) provides a survival advantage over patients who do not achieve this response or achieve it later. In BP, 21% of previously treated patients and 36% of previously untreated patients have responded to imatinib, and up to 17% of patients may achieve a major cytogenetic response. However, responses are frequently short-lived. Several agents are being investigated for treatment of advanced-phase CML, including **decitabine** (DAC), homoharringtonine (HHT), troxacitabine, clofarabine, farnesyl transferase (FTase) inhibitors (FTI), and others. Many have also proven to be synergistic with imatinib in vitro and combination studies are ongoing. Continued investigation of these approaches is needed to improve the long-term prognosis of advanced-phase CML. *Semin Hematol* 40:79-86.

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ACCESSION NUMBER: 2003053879 IN-PROCESS
DOCUMENT NUMBER: 22451283 PubMed ID: 12563614
TITLE: Advanced-phase chronic myeloid leukemia.
AUTHOR: Cortes Jorge; Kantarjian Hagop
CORPORATE SOURCE: Department of Leukemia, The University of Texas, M.D. Anderson Cancer Center, Houston, TX.
SOURCE: SEMINARS IN HEMATOLOGY, (2003 Jan) 40 (1) 79-86.
Journal code: 0404514. ISSN: 0037-1963.
PUB. COUNTRY: United States
DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
LANGUAGE: English
FILE SEGMENT: IN-PROCESS; NONINDEXED; Priority Journals
ENTRY DATE: Entered STN: 20030204
Last Updated on STN: 20030204

L4 ANSWER 2 OF 7 MEDLINE on STN

AB The treatment options for chronic myelogenous leukemia (CML) continue to evolve rapidly. **Imatinib mesylate** (Gleevec, Glivec, formerly ST1571) has continued to show remarkable clinical benefits and the updated results with this agent are reviewed. As relapses using single agent imatinib have occurred, particularly in advanced phase patients, the issue of whether combinations of other antileukemic agents with imatinib may yield improved results is addressed. In addition, data on new agents that have potential in the treatment of CML are reviewed. These agents are presented in the context of their molecular mechanism of action. The most recent data for stem cell transplantation, along with advances in nonmyeloablative transplants, are also reviewed. In Section I, Drs. Stephen O'Brien and Brian Druker update the current status of clinical trials with imatinib and review ongoing investigations into mechanisms of resistance and combinations of imatinib with other agents. They also present their views on integration of imatinib with other therapies. In Section II, Dr. Jorge Cortes describes the most recent data on novel therapies for CML, including farnesyl transferase inhibitors, arsenic trioxide, **decitabine**, and troxatyl, among others. These agents are discussed in the context of their molecular mechanism of action and rationale for use. In Section III, Dr. Jerald Radich updates the results of stem cell transplants for CML, including emerging data on nonmyeloablative transplants. He also presents data on using microarrays to stratify patients into molecularly defined risk groups.

ACCESSION NUMBER: 2002687859 IN-PROCESS
DOCUMENT NUMBER: 22335953 PubMed ID: 12446421
TITLE: Chronic myelogenous leukemia.
AUTHOR: Druker Brian J; O'Brien Stephen G; Cortes Jorge; Radich Jerald
CORPORATE SOURCE: University of Newcastle, Royal Victoria Infirmary, Newcastle Upon Tyne, United Kingdom.
SOURCE: Hematology (Am Soc Hematol Educ Program), (2002) 111-35.

Journal code: 100890099. ISSN: 1520-4391.

PUB. COUNTRY:

United States

DOCUMENT TYPE:

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE:

English

FILE SEGMENT:

IN-PROCESS; NONINDEXED; Priority Journals

ENTRY DATE:

Entered STN: 20021214

Last Updated on STN: 20030713

L4 ANSWER 3 OF 7 MEDLINE on STN

AB Chronic myelogenous leukemia (CML) is a clonal myeloproliferative disorder molecularly defined by the BCR-ABL gene and its products. The protein encoded by this chimeric gene is a constitutively activated tyrosine kinase that alters multiple signal transduction pathways inducing malignant transformation. Until recently, treatment options for patients with CML consisted of hydroxyurea, interferon-based therapies or allogeneic stem cell transplantation (alloSCT). Treatment decisions were generally based on the age of the patient and the phase of the disease. Recently, several new therapies have been developed that may change the natural history of CML and patient prognosis. In particular **imatinib mesylate** (ST1571, Gleevec) an oral Bcr-Abl kinase inhibitor, has demonstrated activity in all phases of CML, and may replace interferon and alloSCT as the initial therapy for this disease. Other agents and therapies with potential value, either alone or in combination, include polyethyleneglycol (PEG) interferon, homoharringtonine, **decitabine**, oral cytarabine, and growth factor modulation. In this article, we discuss the biological and clinical characteristics of CML, as well as the different therapeutic alternatives for patients with this disorder.

ACCESSION NUMBER: 2002254399 MEDLINE

DOCUMENT NUMBER: 21989084 PubMed ID: 11993784

TITLE: Current therapy of chronic myelogenous leukemia.

AUTHOR: Garcia-Manero Guillermo; Talpaz Moshe; Kantarjian Hagop M

CORPORATE SOURCE: Department of Leukemia and Bioimmunotherapy, University of Texas M.D. Anderson Cancer Center, Houston 77030, USA.

SOURCE: INTERNAL MEDICINE, (2002 Apr) 41 (4) 254-64. Ref: 81
Journal code: 9204241. ISSN: 0918-2918.

PUB. COUNTRY: Japan

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

General Review; (REVIEW)
(REVIEW, TUTORIAL)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200211

ENTRY DATE: Entered STN: 20020508

Last Updated on STN: 20021211

Entered Medline: 20021104

L4 ANSWER 4 OF 7 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

AB Clinical phase I/II studies with the Abl kinase inhibitor **imatinib mesylate** (Gleevec/Glivec, formerly ST1571) for the treatment for chronic myelogenous leukemia (CML) demonstrated the safety and the remarkable efficacy of this molecularly targeted agent. However, a significant proportion of patients treated in the chronic phase of the disease after having failed interferon alpha (IFN) remain predominantly Philadelphia chromosome positive (Ph+), suggesting a risk of later relapses. Furthermore, results in blast crisis patients revealed a high frequency of relapses or resistance to imatinib. To circumvent resistance, improve response rates, or prolong survival, pre-clinical evaluations of combinations of imatinib with other agents have been pursued. Some of these have already been translated into clinical studies. Here, we first summarize evidence from pre-clinical studies on new combination regimens with imatinib in the treatment of CML. Second, we analyze preliminary clinical data of ongoing combination studies. Finally, we provide a summary of approaches that use novel antileukemic agents with molecularly characterized modes of action.

ACCESSION NUMBER: 2002:478536 BIOSIS
DOCUMENT NUMBER: PREV200200478536
TITLE: Insights from pre-clinical studies for new combination treatment regimens with the Bcr-Abl kinase inhibitor **imatinib mesylate** (Gleevec/Glivec) in chronic myelogenous leukemia: A translational perspective.
AUTHOR(S): La Rosee, P.; O'Dwyer, M. E.; Druker, B. J. (1)
CORPORATE SOURCE: (1) Division of Hematology and Medical Oncology, Oregon Health and Science University, 3181 Sam Jackson Park Rd, Mail Code L592, Portland, OR, 97201 USA
SOURCE: Leukemia (Basingstoke), (July, 2002) Vol. 16, No. 7, pp. 1213-1219. <http://www.naturesj.com/leu/index.html>. print.
ISSN: 0887-6924.
DOCUMENT TYPE: General Review
LANGUAGE: English

L4 ANSWER 5 OF 7 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AB Chronic myelogenous leukemia (CML) is a clonal myeloproliferative disorder molecularly defined by the BCR-ABL gene and its products. The protein encoded by this chimeric gene is a constitutively activated tyrosine kinase that alters multiple signal transduction pathways inducing malignant transformation. Until recently, treatment options for patients with CML consisted of hydroxyurea, interferon-based therapies or allogeneic stem cell transplantation (alloSCT). Treatment decisions were generally based on the age of the patient and the phase of the disease. Recently, several new therapies have been developed that may change the natural history of CML and patient prognosis. In particular **imatinib mesylate** (ST1571, Gleevec) an oral Bcr-Abl kinase inhibitor, has demonstrated activity in all phases of CML, and may replace interferon and alloSCT as the initial therapy for this disease. Other agents and therapies with potential value, either alone or in combination, include polyethyleneglycol (PEG) interferon, homoharringtonine, **decitabine**, oral cytarabine, and growth factor modulation. In this article, we discuss the biological and clinical characteristics of CML, as well as the different therapeutic alternatives for patients with this disorder.

ACCESSION NUMBER: 2002:376988 BIOSIS
DOCUMENT NUMBER: PREV200200376988
TITLE: Current therapy of chronic myelogenous leukemia.
AUTHOR(S): Garcia-Manero, Guillermo; Talpaz, Moshe; Kantarjian, Hagop M. (1)
CORPORATE SOURCE: (1) Department of Leukemia, University of Texas M. D. Anderson Cancer Center, 1515 Holcombe Blvd, Box 428, Houston, TX, 77030 USA
SOURCE: Internal Medicine (Tokyo), (April, 2002) Vol. 41, No. 4, pp. 254-264. print.
ISSN: 0918-2918.
DOCUMENT TYPE: General Review
LANGUAGE: English

L4 ANSWER 6 OF 7 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AB 237 adult patients (pts) with Ph+ CML AP were treated with **imatinib mesylate** 400-600 mg P.O. daily at our institution as part of 2 Novartis sponsored multi-institutional multinational studies: Novartis 109 the pivotal study (N=58) and Novartis 114 the expanded access study (N=179). 193 pts are evaluable with more than 3 months of follow-up. 156 pts had the classical CML AP criteria (Cancer 61:1441, 1988); 33 pts were treated for blasts 10-14%, blasts+pros 20-29%, or spleen \geq 10 cm \geq 50% increase over 4 weeks (modified CML-AP criteria); 4 pts had second chronic phase. 26 received **imatinib mesylate** 400 mg/D, and 167 pts had **imatinib mesylate** 600 mg/D. Their median age was 50 years. Overall, 162 pts (84%) achieved CHR, 107 (55%) had a cytogenetic response (Ph<90%): major (Ph<35%) in 79 (41%); complete (Ph 0%) in 57

(30%). With a median follow up of 8.4 months, 167 patients (87%) are alive. The estimated 1.5-year survival rate was 75%, and remission duration rate 61%. Prognostic factors associated with lower major CG response rates (pgtoreq0.02) were: age gtoreq60 yrs, splenomegaly gtoreq10 cm bcm, longer duration of chronic phase >3 yrs, WBC >10X10⁹/L, marrow blasts gtoreq15%, and STI dose 400 mg daily. Prognostic factors associated with worse survival (p<0.02) were: age gtoreq60 yrs, hemoglobin <10 g/dl marrow blasts gtoreq15%, cytogenetic clonal evolution and STI dose 400 mg daily and failure to achieve major CG response. Patients treated with 600 vs 400 mg had significantly better major (44% vs 19%, p=0.02) and complete (32% vs 15%, p=0.11) CG response rates, and 1.5 yr survival rates (78% vs 67%, p<0.01). Patients with "modified" CML AP criteria had similar major CG response and survival rates. By multivariate analysis, factors independently predictive negatively for major CG response were (p<0.05): diagnosis to therapy >3 years, and spleen size >10 cm bcm. Those associated with worse survival were (p<0.05): older age, failure to achieve major cytogenetic response, and cytogenetic clonal evolution. In summary **imatinib mesylate** is the most active single agent therapy in accelerated phase. **Imatinib mesylate** combinations with interferon alpha, cytarabine, homoharringtonine, **decitabine** or others are warranted in CML AP.

ACCESSION NUMBER: 2002:153049 BIOSIS
DOCUMENT NUMBER: PREV200200153049
TITLE: Treatment of accelerated phase of Philadelphia chromosome positive chronic myeloid leukemia (Ph+ CML AP) with **imatinib mesylate** (STI571).
AUTHOR(S): Kantarjian, Hagop M. (1); O'Brien, Susan (1); Cortes, Jorge (1); Faderl, Stefan (1); Giles, Francis (1); Thomas, Deborah (1); Garcia-Manero, Guillermo (1); Albitar, Maher; Rios, Mary Beth (1); Shan, Jenny (1); Issa, Jean-Pierre (1); Resta, Debra; Capdeville, Renaud; Keating, Michael J. (1); Freireich, Emil J. (1); Talpaz, Moshe (1)
CORPORATE SOURCE: Leukemia, University of Texas M.D. Anderson Cancer Center, Houston, TX USA
SOURCE: Blood, (November 16, 2001) Vol. 98, No. 11 Part 1, pp. 141a. <http://www.bloodjournal.org/>. print.
Meeting Info.: 43rd Annual Meeting of the American Society of Hematology, Part 1 Orlando, Florida, USA December 07-11, 2001
ISSN: 0006-4971.
DOCUMENT TYPE: Conference
LANGUAGE: English

L4 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2003 ACS on STN
AB Methods, compns. and kits are provided for treating cancer assocd. with protein tyrosine kinase activity such as chronic myelogenous leukemia. In particular, a treatment method is provided comprising: administering to a patient having chronic myelogenous leukemia and a degree of resistance to **imatinib mesylate**, a therapeutically effective amt. of a DNA methylation inhibitor which mitigates the **imatinib mesylate** resistance.

ACCESSION NUMBER: 2003:609844 CAPLUS
TITLE: Method for treating chronic myelogenous leukemia combined with some resistance to **imatinib mesylate** using DNA methylation inhibitor to mitigate **imatinib mesylate** resistance
INVENTOR(S): Lyons, John
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 10 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003147813	A1	20030807	US 2002-71849	20020207
WO 2003065995	A2	20030814	WO 2003-US3537	20030206
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			US 2002-71849	A1 20020207
			US 2002-206854	A1 20020726

=> s 12 and treatment?

L5 436 L2 AND TREATMENT?

=> s 12 and dna(w)methylation?

L6 3 L2 AND DNA(W) METHYLATION?

=> d 16 abs ibib 1-3

L6 ANSWER 1 OF 3 MEDLINE on STN

AB Very promising results have been obtained in clinical trials on chronic-phase chronic myeloid leukemia (CP-CML) patients treated with **imatinib mesylate** (IM; Gleevecr, ST1571), a BCR-ABL tyrosine kinase inhibitor. However, we found that IM caused considerable inhibition of normal hematopoietic progenitor cells upon treating control bone marrow (BM) cultures. In vitro IM treatment gave a decrease in the yield and size of colonies from BM of untreated CP-CML patients that was only two to three times that from the normal samples. Moreover, about 30% of myeloid progenitors (CFU-GM) from CML BM still formed colonies in the presence of IM, most of which had BCR-ABL RNA. About half of these treated colonies also displayed methylation of the internal ABL Pa promoter, a CML-specific epigenetic alteration, which was used in this study as a marker for BCR-ABL translocation-containing cells. However, 5-8% of the treated or the untreated CML BM-derived colonies had no detectable BCR-ABL RNA by two or three rounds of RT-PCR despite being positive for the internal standard RNA and displaying hallmarks of CML, either t(9;22)(q34;ql 1) or ABL Pa methylation. Our results indicate that IM is only partially specific for CML progenitor cells compared to normal hematopoietic progenitor cells and suggest that some CML cells may have a silent BCR-ABL oncogene that could interfere with therapy.

ACCESSION NUMBER: 2003156874 MEDLINE

DOCUMENT NUMBER: 22560189 PubMed ID: 12673129

TITLE: Imatinib (ST1571) provides only limited selectivity for CML cells and treatment might be complicated by silent BCR-ABL genes.

COMMENT: Comment in: Cancer Biol Ther. 2003 Jan-Feb;2(1):109-10

AUTHOR: Jiang Guanchao; Yang Fan; Li Marilyn; Weissbecker Karen; Price Sherrie; Kim K C; La Russa Vincent F; Safah Hana; Ehrlich Melanie

CORPORATE SOURCE: Tulane Cancer Center and Human Genetics Program, Tulane Medical School, New Orleans, Louisiana 70112, USA.

CONTRACT NUMBER: CA78639 (NCI)

SOURCE: Cancer Biol Ther, (2003 Jan-Feb) 2 (1) 103-8.

Journal code: 101137842. ISSN: 1538-4047.

PUB. COUNTRY: United States
 DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
 LANGUAGE: English
 FILE SEGMENT: Priority Journals
 ENTRY MONTH: 200307
 ENTRY DATE: Entered STN: 20030404
 Last Updated on STN: 20030724
 Entered Medline: 20030723

L6 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN
 AB Methods are provided for treating diseases associated with abnormal activity of kinases such as chronic myelogenous leukemia. The method comprises: administering a **DNA methylation inhibitor** to the patient in therapeutically effective amount; and administering a kinase inhibitor such as **imatinib mesylate** to the patient in therapeutically effective amount, such that the *in vivo* activity of the kinase is reduced relative to that prior to the treatment. The method can be used to treat cancer associated with abnormal activity of kinases such as phosphatidylinositol 3'-kinase (P13K), protein kinases including serine/threonine kinases such as Raf kinases, protein kinase kinases such as MEK, and tyrosine kinases such as those in the epidermal growth factor receptor family (EGFR), platelet-derived growth factor receptor family (PDGFR), vascular endothelial growth factor receptor (VEGFR) family, nerve growth factor receptor family (NGFR), fibroblast growth factor receptor family (FGFR) insulin receptor family, ephrin receptor family, Met family, Ror family, c-kit family, Src family, Fes family, JAK family, Fak family, Btk family, Syk/ZAP-70 family, and Abl family.

ACCESSION NUMBER: 2003:633416 CAPLUS
 TITLE: Method for treating diseases associated with abnormal kinase activity
 INVENTOR(S): Lyons, John; Rubinfeld, Joseph
 PATENT ASSIGNEE(S): Supergen, Inc., USA
 SOURCE: PCT Int. Appl., 64 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003065995	A2	20030814	WO 2003-US3537	20030206
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
US 2003147813	A1	20030807	US 2002-71849	20020207
PRIORITY APPLN. INFO.:			US 2002-71849	A1 20020207
			US 2002-206854	A1 20020726

L6 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN
 AB Methods, compns. and kits are provided for treating cancer assocd. with protein tyrosine kinase activity such as chronic myelogenous leukemia. In particular, a treatment method is provided comprising: administering to a patient having chronic myelogenous leukemia and a degree of resistance to **imatinib mesylate**, a therapeutically effective amt. of a **DNA**

methylation inhibitor which mitigates the imatinib mesylate resistance.

ACCESSION NUMBER: 2003:609844 CAPLUS
TITLE: Method for treating chronic myelogenous leukemia combined with some resistance to imatinib mesylate using DNA methylation inhibitor to mitigate imatinib mesylate resistance
INVENTOR(S): Lyons, John
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 10 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003147813	A1	20030807	US 2002-71849	20020207
WO 2003065995	A2	20030814	WO 2003-US3537	20030206
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			US 2002-71849	A1 20020207
			US 2002-206854	A1 20020726

=> s dna(w)methylation and l1
L7 3 DNA(W) METHYLATION AND L1

=> d 17 1-3

L7 ANSWER 1 OF 3 MEDLINE on STN
AN 2003156874 MEDLINE
DN 22560189 PubMed ID: 12673129
TI Imatinib (ST1571) provides only limited selectivity for CML cells and treatment might be complicated by silent BCR-ABL genes.
CM Comment in: Cancer Biol Ther. 2003 Jan-Feb;2(1):109-10
AU Jiang Guanchao; Yang Fan; Li Marilyn; Weissbecker Karen; Price Sherrie; Kim K C; La Russa Vincent F; Safah Hana; Ehrlich Melanie
CS Tulane Cancer Center and Human Genetics Program, Tulane Medical School, New Orleans, Louisiana 70112, USA.
NC CA78639 (NCI)
CA81506 (NCI)
SO Cancer Biol Ther, (2003 Jan-Feb) 2 (1) 103-8.
Journal code: 101137842. ISSN: 1538-4047.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200307
ED Entered STN: 20030404
Last Updated on STN: 20030724
Entered Medline: 20030723

L7 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

AN 2003:633416 CAPLUS
TI Method for treating diseases associated with abnormal kinase activity
IN Lyons, John; Rubinfeld, Joseph
PA Supergen, Inc., USA
SO PCT Int. Appl., 64 pp.
CODEN: PIXXD2

DT Patent
LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003065995	A2	20030814	WO 2003-US3537	20030206
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	US 2003147813	A1	20030807	US 2002-71849	20020207
PRAI	US 2002-71849	A1	20020207		
	US 2002-206854	A1	20020726		

L7 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

AN 2003:609844 CAPLUS

TI Method for treating chronic myelogenous leukemia combined with some resistance to imatinib mesylate using DNA methylation inhibitor to mitigate imatinib mesylate resistance

IN Lyons, John

PA USA

SO U.S. Pat. Appl. Publ., 10 pp.
CODEN: USXXCO

DT Patent
LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003147813	A1	20030807	US 2002-71849	20020207
	WO 2003065995	A2	20030814	WO 2003-US3537	20030206
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI	US 2002-71849	A1	20020207		
	US 2002-206854	A1	20020726		

=> s dna(w)methylation and inhibitor?

L8 2446 DNA(W) METHYLATION AND INHIBITOR?

=> s l8 and l1

L9 3 L8 AND L1

=> s 19 1-3

MISSING OPERATOR L9 1-3

The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> d 19 1-3

L9 ANSWER 1 OF 3 MEDLINE on STN
AN 2003156874 MEDLINE
DN 22560189 PubMed ID: 12673129
TI Imatinib (ST1571) provides only limited selectivity for CML cells and treatment might be complicated by silent BCR-ABL genes.
CM Comment in: Cancer Biol Ther. 2003 Jan-Feb;2(1):109-10
AU Jiang Guanchao; Yang Fan; Li Marilyn; Weissbecker Karen; Price Sherrie; Kim K C; La Russa Vincent F; Safah Hana; Ehrlich Melanie
CS Tulane Cancer Center and Human Genetics Program, Tulane Medical School, New Orleans, Louisiana 70112, USA.
NC CA78639 (NCI)
CA81506 (NCI)
SO Cancer Biol Ther, (2003 Jan-Feb) 2 (1) 103-8.
Journal code: 101137842. ISSN: 1538-4047.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200307
ED Entered STN: 20030404
Last Updated on STN: 20030724
Entered Medline: 20030723

L9 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN
AN 2003:633416 CAPLUS
TI Method for treating diseases associated with abnormal kinase activity
IN Lyons, John; Rubinfeld, Joseph
PA Supergen, Inc., USA
SO PCT Int. Appl., 64 pp.
CODEN: PIXXD2

DT Patent
LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003065995	A2	20030814	WO 2003-US3537	20030206
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI	US 2003147813	A1	20030807	US 2002-71849	20020207
	US 2002-71849	A1	20020207		
	US 2002-206854	A1	20020726		

L9 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

AN 2003:609844 CAPLUS

TI Method for treating chronic myelogenous leukemia combined with some resistance to imatinib mesylate using DNA methylation inhibitor to mitigate imatinib mesylate resistance
IN Lyons, John

PA USA
SO U.S. Pat. Appl. Publ., 10 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 2003147813	A1	20030807	US 2002-71849	20020207
	WO 2003065995	A2	20030814	WO 2003-US3537	20030206
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRAI	US 2002-71849	A1	20020207		
	US 2002-206854	A1	20020726		

=> s myelogenous (w) leukemia and staged?
L10 3 MYELOGENOUS (W) LEUKEMIA AND STAGED?

=> s myelogenous (w) leukemia?
L11 24614 MYELOGENOUS (W) LEUKEMIA?

=> s 111 and cytidine?
L12 70 L11 AND CYTIDINE?

=> d 112 1-70

L12 ANSWER 1 OF 70 MEDLINE on STN
AN 2002222196 MEDLINE
DN 21956901 PubMed ID: 11960335
TI Results of a phase II trial of a combination of oral cytarabine ocfosfate (YNK01) and interferon alpha-2b for the treatment of chronic myelogenous leukemia patients in chronic phase.
AU Maloisel F; Guerci A; Guyotat D; Ifrah N; Michallet M; Reiffers J; Tertain G; Blanc M; Bauduer F; Briere J; Abgrall J F; Pegourie-Bandelier B; Solary E; Cambier N; Coso D; Vilque J P; Delain M; Harousseau J L; Rousselot P; Belhadj K; Morice P; Attal J; Chabin M; Chastang C; Guilhot J; Guilhot F
CS Division of Hematology, University Hospital of Strasbourg, France. (France Intergrroupe des Leucemies Myeloides Chroniques).
SO LEUKEMIA, (2002 Apr) 16 (4) 573-80.
CY England: United Kingdom
DT (CLINICAL TRIAL)
(CLINICAL TRIAL, PHASE II)
Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200205
ED Entered STN: 20020418
Last Updated on STN: 20020508
Entered Medline: 20020507

L12 ANSWER 2 OF 70 MEDLINE on STN
AN 2002206547 MEDLINE
DN 21936329 PubMed ID: 11939268
TI Bone marrow cytogenetic complete remission achieved by interferon-alpha

plus cytarabine ocfosfate therapy in a patient with chronic myelogenous leukemia during extramedullary blast crisis.

AU Gotoh Akihiko; Miyazawa Keisuke; Uchida Yoshiko; Sashida Goro; Kawakubo Ken; Kuriyama Yuzuru; Ohyashiki Kazuma
CS First Department of Internal Medicine, Tokyo Medical University, Japan.
SO INTERNATIONAL JOURNAL OF HEMATOLOGY, (2002 Feb) 75 (2) 191-4.
Journal code: 9111627. ISSN: 0925-5710.

CY Ireland
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200301
ED Entered STN: 20020410
Last Updated on STN: 20030125
Entered Medline: 20030124

L12 ANSWER 3 OF 70 MEDLINE on STN
AN 2001238195 MEDLINE
DN 21218123 PubMed ID: 11320667
TI Comparative study of a novel nucleoside analogue (Troxatyl, troxacicabine, BCH-4556) and AraC against leukemic human tumor xenografts expressing high or low cytidine deaminase activity.
AU Gourdeau H; Bibeau L; Ouellet F; Custeau D; Bernier L; Bowlin T
CS BioChem Pharma Inc., 275 Armand-Frappier Blvd, Laval, Quebec H7V 4A7, Canada.. gourdeah@biochempharma.com
SO CANCER CHEMOTHERAPY AND PHARMACOLOGY, (2001 Mar) 47 (3) 236-40.
Journal code: 7806519. ISSN: 0344-5704.
CY Germany: Germany, Federal Republic of
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200105
ED Entered STN: 20010517
Last Updated on STN: 20020420
Entered Medline: 20010503

L12 ANSWER 4 OF 70 MEDLINE on STN
AN 2001100720 MEDLINE
DN 21036706 PubMed ID: 11196156
TI Simultaneous treatment with 1-beta-D-arabinofuranosylcytosine and daunorubicin induces cross-resistance to both drugs due to a combination-specific mechanism in HL60 cells.
AU Takemura H; Urasaki Y; Yoshida A; Fukushima T; Ueda T
CS First Department of Internal Medicine, Fukui Medical University, Matsuoka, Japan.
SO CANCER RESEARCH, (2001 Jan 1) 61 (1) 172-7.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200102
ED Entered STN: 20010322
Last Updated on STN: 20010322
Entered Medline: 20010201

L12 ANSWER 5 OF 70 MEDLINE on STN
AN 2000421721 MEDLINE
DN 20327793 PubMed ID: 10867132
TI Treatment of patients with advanced chronic myelogenous leukemia with interferon-alpha-2b and continuous oral cytarabine ocfosfate (YNK01): a pilot study.
AU Kuhr T; Eisterer W; Apfelbeck U; Linkesch W; Bechter O; Zabernigg A; Geissler K; Barbieri G; Duba C; Gastl G; Thaler J
CS Department of Internal Medicine, University Hospital, Anichstrasse 35,

SO 6020, Innsbruck, Austria.. thomas.kuehr@uibk.ac.at
LEUKEMIA RESEARCH, (2000 Jul) 24 (7) 583-7.
Journal code: 7706787. ISSN: 0145-2126.

CY ENGLAND: United Kingdom
DT (CLINICAL TRIAL)
Journal; Article; (JOURNAL ARTICLE)
(MULTICENTER STUDY)

LA English
FS Priority Journals
EM 200009
ED Entered STN: 20000915
Last Updated on STN: 20000915
Entered Medline: 20000907

L12 ANSWER 6 OF 70 MEDLINE on STN
AN 2000084096 MEDLINE
DN 20084096 PubMed ID: 10616723
TI Isolation and characterization of 5-carbamoylmethyluridine and 5-carbamoylmethyl-2-thiouridine from human urine.
AU Chheda G B; Patrzyc H B; Tworek H A; Dutta S P
CS Department of Biophysics, Roswell Park Cancer Institute, Buffalo, NY 14263, USA.
SO NUCLEOSIDES AND NUCLEOTIDES, (1999 Oct) 18 (10) 2155-73.
Journal code: 8215930. ISSN: 0732-8311.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 200002
ED Entered STN: 20000229
Last Updated on STN: 20000229
Entered Medline: 20000214

L12 ANSWER 7 OF 70 MEDLINE on STN
AN 1998339469 MEDLINE
DN 98339469 PubMed ID: 9676847
TI Accumulation of arabinosyluracil 5'-triphosphate during arabinosylcytosine therapy in circulating blasts of patients with acute **myelogenous** leukemia.
AU Gandhi V; Xu Y Z; Estey E
CS Department of Clinical Investigation, The University of Texas M.D. Anderson Cancer Center, Houston 77030, USA.
NC CA32839 (NCI)
CA55164 (NCI)
CA57629 (NCI)
SO CLINICAL CANCER RESEARCH, (1998 Jul) 4 (7) 1719-26.
Journal code: 9502500. ISSN: 1078-0432.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199809
ED Entered STN: 19980925
Last Updated on STN: 19980925
Entered Medline: 19980916

L12 ANSWER 8 OF 70 MEDLINE on STN
AN 1998240988 MEDLINE
DN 98240988 PubMed ID: 9581832
TI Telomerase from human leukemia cells: properties and its interaction with deoxynucleoside analogues.
AU Pai R B; Pai S B; Kukhanova M; Dutschman G E; Guo X; Cheng Y C
CS Department of Pharmacology, Yale School of Medicine, Yale University, New Haven, Connecticut 06510, USA.
NC AI-38204 (NIAID)

SO CANCER RESEARCH, (1998 May 1) 58 (9) 1909-13.
Journal code: 2984705R. ISSN: 0008-5472.

CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199806
ED Entered STN: 19980611
Last Updated on STN: 19980611
Entered Medline: 19980602

L12 ANSWER 9 OF 70 MEDLINE on STN
AN 97083073 MEDLINE
DN 97083073 PubMed ID: 8929647
TI Combination therapy with granulocyte colony-stimulating factor, all-trans retinoic acid, and low-dose cytotoxic drugs for acute **myelogenous leukemia**.
AU Usuki K; Kitazume K; Endo M; Ito K; Iki S; Urabe A
CS Division of Hematology, Kanto Teishin Hospital, Tokyo.
SO INTERNAL MEDICINE, (1995 Dec) 34 (12). 1186-9.
Journal code: 9204241. ISSN: 0918-2918.
CY Japan
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199703
ED Entered STN: 19970407
Last Updated on STN: 19970407
Entered Medline: 19970324

L12 ANSWER 10 OF 70 MEDLINE on STN
AN 95275050 MEDLINE
DN 95275050 PubMed ID: 7755392
TI Low-dose cytarabine ocfosfate therapy in an elderly acute **myelogenous leukemia**.
AU Hamaoka R; Jozaki K; Amano T; Itoh H; Imai Y; Nishikawa M; Kurokawa M; Yonezawa T; Chinen Y
CS Dept. of Internal Medicine, Ikeda Municipal Hospital.
SO GAN TO KAGAKU RYOH [JAPANESE JOURNAL OF CANCER AND CHEMOTHERAPY], (1995 May) 22 (6) 819-22.
Journal code: 7810034. ISSN: 0385-0684.
CY Japan
DT Journal; Article; (JOURNAL ARTICLE)
LA Japanese
FS Priority Journals
EM 199506
ED Entered STN: 19950629
Last Updated on STN: 19950629
Entered Medline: 19950616

L12 ANSWER 11 OF 70 MEDLINE on STN
AN 94175542 MEDLINE
DN 94175542 PubMed ID: 8129396
TI Successful treatment of acute **myelogenous leukemia** in an elderly patient with cytarabine ocfosfate.
AU Inaba T; Shimazaki C; Tatsumi T; Yamagata N; Hirata T; Goto H; Fujita N; Nakagawa M; Fujita N; Miyazaki S; +
CS Second Dept. of Medicine, Kyoto Prefectural University of Medicine.
SO GAN TO KAGAKU RYOH [JAPANESE JOURNAL OF CANCER AND CHEMOTHERAPY], (1994 Mar) 21 (4) 535-8.
Journal code: 7810034. ISSN: 0385-0684.
CY Japan
DT Journal; Article; (JOURNAL ARTICLE)
LA Japanese
FS Priority Journals

EM 199404
ED Entered STN: 19940420
Last Updated on STN: 19940420
Entered Medline: 19940412

L12 ANSWER 12 OF 70 MEDLINE on STN
AN 94034807 MEDLINE
DN 94034807 PubMed ID: 8220157
TI Role of aberrant sialylation of chronic myeloid leukemia granulocytes on binding and signal transduction by chemotactic peptides and colony stimulating factors.
AU Cyopick P; Culliton R; Brockhausen I; Sutherland D R; Mills G B; Baker M
CS Toronto Hospital, Ontario, Canada.
SO LEUKEMIA AND LYMPHOMA, (1993 Sep) 11 (1-2) 79-90.
Journal code: 9007422. ISSN: 1042-8194.
CY Switzerland
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199312
ED Entered STN: 19940117
Last Updated on STN: 19970203
Entered Medline: 19931207

L12 ANSWER 13 OF 70 MEDLINE on STN
AN 91339133 MEDLINE
DN 91339133 PubMed ID: 1873797
TI Hemin enhances the sensitivity of erythroleukemia cells to 1-beta-D-arabinofuranosylcytosine by both activation of deoxycytidine kinase and reduction of cytidine deaminase activity.
AU Honma Y; Onozuka Y; Okabe-Kado J; Kasukabe T; Hozumi M
CS Department of Chemotherapy, Saitama Cancer Center Research Institute, Japan.
SO CANCER RESEARCH, (1991 Sep 1) 51 (17) 4535-8.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199109
ED Entered STN: 19911013
Last Updated on STN: 19980206
Entered Medline: 19910923

L12 ANSWER 14 OF 70 MEDLINE on STN
AN 91199087 MEDLINE
DN 91199087 PubMed ID: 1707752
TI Effects of 2-chloro-9-(2-deoxy-2-fluoro-beta-D-arabinofuranosyl)adenine on K562 cellular metabolism and the inhibition of human ribonucleotide reductase and DNA polymerases by its 5'-triphosphate.
AU Parker W B; Shaddix S C; Chang C H; White E L; Rose L M; Brockman R W; Shortnacy A T; Montgomery J A; Secrist J A 3rd; Bennett L L Jr
CS Kettering-Meyer Laboratory, Southern Research Institute, Birmingham, Alabama 35205.
NC CA34200 (NCI)
SO CANCER RESEARCH, (1991 May 1) 51 (9) 2386-94.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199105
ED Entered STN: 19910607
Last Updated on STN: 19980206
Entered Medline: 19910517

L12 ANSWER 15 OF 70 MEDLINE on STN
AN 91004045 MEDLINE
DN 91004045 PubMed ID: 2208147
TI Pharmacologically directed design of the dose rate and schedule of 2',2'-difluorodeoxycytidine (Gemcitabine) administration in leukemia.
AU Grunewald R; Kantarjian H; Keating M J; Abbruzzese J; Tarassoff P; Plunkett W
CS Department of Medical Oncology, University of Texas, M.D. Anderson Cancer Center, Houston 77030.
NC CA32839 (NCI)
SO CANCER RESEARCH, (1990 Nov 1) 50 (21) 6823-6.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199011
ED Entered STN: 19910117
Last Updated on STN: 19910117
Entered Medline: 19901121

L12 ANSWER 16 OF 70 MEDLINE on STN
AN 90335802 MEDLINE
DN 90335802 PubMed ID: 2379165
TI Human leukemic myeloblasts and myeloblastoid cells contain the enzyme cytidine 5'-monophosphate-N-acetylneurameric acid:Gal beta 1-3GalNAc alpha (2-3)-sialyltransferase.
AU Kanani A; Sutherland D R; Fibach E; Matta K L; Hindenburg A; Brockhausen I; Kuhns W; Taub R N; van den Eijnden D H; Baker M A
CS Department of Medicine, Toronto General Hospital, Ontario, Canada.
NC CA31762 (NCI)
CA35329 (NCI)
SO CANCER RESEARCH, (1990 Aug 15) 50 (16) 5003-7.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 199009
ED Entered STN: 19901012
Last Updated on STN: 19980206
Entered Medline: 19900912

L12 ANSWER 17 OF 70 MEDLINE on STN
AN 88310800 MEDLINE
DN 88310800 PubMed ID: 2457428
TI Effect of HpaII and MspI restriction endonucleases on chronic myelogenous leukemia chromosomes. Detection of CpG dinucleotide demethylation in situ.
AU Ferrucci L; Mezzanotte R; Vanni R; Stuppia R; Guanciali-Franchi P; Calabrese G; Palka G; Bianchi U; Sumner A T
CS Dipartimento di Biologia, Facolta di Science M.F.N., II Universita di Roma, Italy.
SO CANCER GENETICS AND CYTOGENETICS, (1988 Sep) 34 (2) 251-6.
Journal code: 7909240. ISSN: 0165-4608.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 198810
ED Entered STN: 19900308
Last Updated on STN: 19960129
Entered Medline: 19881007

L12 ANSWER 18 OF 70 MEDLINE on STN
AN 87187152 MEDLINE
DN 87187152 PubMed ID: 3471317
TI Presence of cytidine 5'-monophospho-N-acetylneurameric acid:Gal beta 1-3GalNAc-R alpha(2-3)-sialyltransferase in normal human leukocytes and increased activity of this enzyme in granulocytes from chronic myelogenous leukemia patients.
AU Baker M A; Kanani A; Brockhausen I; Schachter H; Hindenburg A; Taub R N
NC CA 31761 (NCI)
CA 37162 (NCI)
SO CANCER RESEARCH, (1987 Jun 1) 47 (11) 2763-6.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 198706
ED Entered STN: 19900303
Last Updated on STN: 19980206
Entered Medline: 19870625

L12 ANSWER 19 OF 70 MEDLINE on STN
AN 82048392 MEDLINE
DN 82048392 PubMed ID: 6945901
TI An in vitro model for acute myelogenous leukemia chemotherapy.
AU Koeffler H P; Yen J; Lowe L
NC CA-15619 (NCI)
CA-15688 (NCI)
CA-16043 (NCI)
+
SO CANCER, (1981 Nov 1) 48 (9) 1958-63.
Journal code: 0374236. ISSN: 0008-543X.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Abridged Index Medicus Journals; Priority Journals
EM 198201
ED Entered STN: 19900316
Last Updated on STN: 19970203
Entered Medline: 19820109

L12 ANSWER 20 OF 70 MEDLINE on STN
AN 78167199 MEDLINE
DN 78167199 PubMed ID: 274175
TI Formation of 1-beta-D-arabinofuranosylcytosine diphosphate choline in neoplastic and normal cells.
AU Lauzon G J; Paterson A R; Belch A W
SO CANCER RESEARCH, (1978 Jun) 38 (6) 1730-3.
Journal code: 2984705R. ISSN: 0008-5472.
CY United States
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 197807
ED Entered STN: 19900314
Last Updated on STN: 19970203
Entered Medline: 19780726

L12 ANSWER 21 OF 70 MEDLINE on STN
AN 76251112 MEDLINE
DN 76251112 PubMed ID: 60073
TI 5-Azacytidine. A new anticancer drug with effectiveness in acute myelogenous leukemia.
AU Von Hoff D D; Slavik M; Muggia F M

SO ANNALS OF INTERNAL MEDICINE, (1976 Aug) 85 (2) 237-45. Ref: 73
Journal code: 0372351. ISSN: 0003-4819.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)
General Review; (REVIEW)

LA English

FS Abridged Index Medicus Journals; Priority Journals

EM 197609

ED Entered STN: 19900313
Last Updated on STN: 19970203
Entered Medline: 19760925

L12 ANSWER 22 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 2001:90082 BIOSIS
DN PREV200100090082
TI Simultaneous treatment with 1-beta-D-arabinofuranosylcytosine and daunorubicin induces cross-resistance to both drugs due to a combination-specific mechanism in HL60 cells.
AU Takemura, Haruyuki; Urasaki, Yoshimasa; Yoshida, Akira; Fukushima, Toshihiro; Ueda, Takanori (1)
CS (1) First Department of Internal Medicine, Fukui Medical University, 23-3, Shimoaizuki, Matsuoka, Fukui, 910-1193 Japan
SO Cancer Research, (January 1, 2001) Vol. 61, No. 1, pp. 172-177. print.
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DT Article
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SL English

L12 ANSWER 23 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
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CS (1) Fukui Med Univ, Fujui Japan
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ISSN: 0197-016X.
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L12 ANSWER 24 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 2000:214465 BIOSIS
DN PREV200000214465
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AU Higuchi, M. (1); Azuma, A.; Matsuda, A.; Sasaki, T.; Fukushima, M.
CS (1) Hokkaido Univ, Sapporo Japan
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Meeting Info.: 91st Annual Meeting of the American Association for Cancer Research. San Francisco, California, USA April 01-05, 2000
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L12 ANSWER 25 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
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DN PREV200000198019

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AU Schwartz, Joseph (1); Alster, Yair; Ben-Tal, Ofira; Lowenstein, Anat
CS (1) Department of Hematology, Tel-Aviv Sourasky Medical Center, 6 Weizman
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DT Article; Letter
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L12 ANSWER 26 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1998:274149 BIOSIS
DN PREV199800274149
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CS (1) Dep. Pharmacol., Yale Sch. Med., Yale Univ., 333 Cedar St., New Haven,
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ISSN: 0008-5472.
DT Article
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L12 ANSWER 27 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1991:457552 BIOSIS
DN BA92:102332
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ARABINOFURANOSYLCYTOSINE BY BOTH ACTIVATION OF DEOXYCYTIDINE KINASE AND
REDUCTION OF CYTIDINE DEAMINASE ACTIVITY.
AU HONMA Y; ONOZUKA Y; OKABE-KADO J; KASUKABE T; HOZUMI M
CS DEP. CHEMOTHERAPY, SAITAMA CANCER CENT., RES. INST., INA, SAITAMA-362,
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CODEN: CNREA8. ISSN: 0008-5472.
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L12 ANSWER 28 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1988:506365 BIOSIS
DN BA86:127049
TI EFFECT OF CYTOSINE ARABINOSIDE ON THE HUMAN IMMUNOSYSTEM METABOLISM AND
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AU VILPO J A; VEROMAA T; EEROLA E
CS LAB. MOLECULAR HEMATOLOGY, BIOCENTER, UNIV. OULU, SF-90220 OULU, FINLAND.
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L12 ANSWER 29 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1987:317770 BIOSIS
DN BA84:37277
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GALACTOSYL-BETA-1-3-N-ACETYL-D-GALACTOSAMINE ALPHA-2-3-SIALYLTRANSFERASE
IN NORMAL HUMAN LEUKOCYTES AND INCREASED ACTIVITY OF THIS ENZYME IN
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L12 ANSWER 30 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1984:332606 BIOSIS
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AU VOGLER W R; WINTON E F; GORDON D S; RANEY M R; GO B; MEYER L
CS 718 WOODRUFF MEMORIAL BUILDING, EMORY UNIVERSITY, ATLANTA, GA. 30322.
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AN 1984:114891 BIOSIS
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CODEN: EXHMA6. ISSN: 0301-472X.
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AN 1983:3353 BIOSIS
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AN 1982:304886 BIOSIS

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TI 5 AZA CYTIDINE IN REFRACTORY ACUTE LEUKEMIA.

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CS DIV OF HEMATOL., DEP. OF MED., MAINE MED. CENT., PORTLAND, OREG. MAINE 04102, USA.

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L12 ANSWER 36 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

AN 1982:239196 BIOSIS

DN BA74:11676

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CS BALTIMORE CANCER RES. CENT., 22 S. GREENE ST., BALTIMORE, MD. 21201.

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DN BA73:55570

TI TREATMENT OF PATIENTS OVER 50 YEARS OF AGE WITH ACUTE MYELOGENOUS LEUKEMIA WITH A COMBINATION OF RUBIDAZONE AND CYTOSINE ARABINOSIDE VINCRISTINE AND PREDNISONE.

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CS DEP. OF DEVELOPMENTAL THERAPEUTICS, M. D. ANDERSON HOSPITAL AND TUMOR INSTITUTE, 6723 BERTNER, HOUSTON, TEX. 77030.

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CS UNIV. CALIFORNIA, DEP. MED., CENT. HEALTH SCI., LOS ANGELES, CALIF. 90024.

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L12 ANSWER 39 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN

AN 1982:20836 BIOSIS

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TI PROGNOSTIC FACTORS AFFECTING REMISSION INDUCTION AND DURATION IN ADULT ACUTE MYELOGENOUS LEUKEMIA.

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CS EMORY UNIV., ATLANTA, GEORGIA 3032.
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AN 1981:7098 BIOSIS
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LEUKEMIA WITH 5 AZA CYTIDINE AND VP-16-213 VEPESIDE.
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L12 ANSWER 41 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1980:156809 BIOSIS
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TI EVALUATION OF CYCLO CYTIDINE IN CHILDREN WITH ADVANCED ACUTE
LEUKEMIA AND SOLID TUMORS.
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CALIF. 90031, USA.
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CODEN: CTRRDO. ISSN: 0361-5960.
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AN 1980:110324 BIOSIS
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DN BR19:6196
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CODEN: PPTCBY. ISBN: 0-8391-1317-.
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L12 ANSWER 44 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1978:191327 BIOSIS
DN BA66:3824
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LEUKEMIC BLAST CELLS.
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CS NATL. INST. HEALTH, ROOM 6N119, BUILD. 10, BETHESDA, MD. 20014, USA.
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L12 ANSWER 45 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1978:90222 BIOSIS
DN BR15:33722
TI TREATMENT OF REFRACTORY ACUTE **MYELOGENOUS LEUKEMIA**
WITH 5 AZA CYTIDINE PLUS BETA DEOXY THIO GUANOSINE.
AU OMURA G A
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CODEN: PAACA3. ISSN: 0569-2296.
DT Conference
FS BR; OLD
LA Unavailable

L12 ANSWER 46 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1978:70838 BIOSIS
DN BR15:14338
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SO Cancer Treat. Rep., (1978) 62 (4), 573-574.
CODEN: CTRRDO. ISSN: 0361-5960.
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L12 ANSWER 47 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1978:49726 BIOSIS
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CODEN: CTRRDO. ISSN: 0361-5960.
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L12 ANSWER 48 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1977:6118 BIOSIS
DN BR13:6118
TI CONTINUOUS INTRA VENOUS INFUSION OF CYCLO CYTIDINE IN ADULT
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CODEN: NKGZAE. ISSN: 0001-5806.
DT Conference
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LA Unavailable

L12 ANSWER 49 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1976:238155 BIOSIS
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CODEN: AIMEAS. ISSN: 0003-4819.

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AN 1975:96778 BIOSIS
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TI TREATMENT OF ACUTE LEUKEMIA WITH 5 AZA CYTIDINE NSC-102816.
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M P
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AN 1974:127490 BIOSIS
DN BA57:27190
TI 5 AZA CYTIDINE A NEW ACTIVE AGENT FOR THE TREATMENT OF ACUTE
LEUKEMIA.
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CODEN: BLOOAW. ISSN: 0006-4971.

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LA Unavailable

L12 ANSWER 53 OF 70 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. on STN
AN 1973:61141 BIOSIS
DN BR09:61141
TI 5 AZA CYTIDINE EFFECTIVE TREATMENT FOR ACUTE LEUKEMIA IN
CHILDREN.
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SO Proc. Am. Assoc. Cancer Res., (1973) 14, 94.
CODEN: PAACAA3. ISSN: 0569-2296.

DT Conference
FS BR; OLD
LA Unavailable

L12 ANSWER 54 OF 70 CAPLUS COPYRIGHT 2003 ACS on STN
AN 2003:609844 CAPLUS
TI Method for treating chronic *myelogenous* leukemia
combined with some resistance to imatinib mesylate using DNA methylation
inhibitor to mitigate imatinib mesylate resistance
IN Lyons, John
PA USA
SO U.S. Pat. Appl. Publ., 10 pp.
CODEN: USXXCO
DT Patent
LA English
FAN.CNT 2

PATENT NO.		KIND	DATE	APPLICATION NO.	DATE
PI	US 2003147813	A1	20030807	US 2002-71849	20020207
	WO 2003065995	A2	20030814	WO 2003-US3537	20030206
				W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM	
				RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG	
PRAI	US 2002-71849	A1	20020207		
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L12	ANSWER 55 OF 70	CAPLUS	COPYRIGHT 2003 ACS on STN		
AN	2001:606409	CAPLUS			
DN	136:303672				
TI	Relationship between clinical efficacy and intracellular levels of dCK and CDA in acute leukemia				
AU	Chen, Fangyuan; Lu, Hongmin; Xuan, Zhenghau; Han, Jieying; Teng, Ye; Ouyang, Renrong				
CS	Department of Hematology, Renji Hospital, Shanghai Second Medical University, Shanghai, 200001, Peop. Rep. China				
SO	Shanghai Yixue (2001), 24(5), 266-269 CODEN: SIHSD8; ISSN: 0253-9934				
PB	Shanghai Yixue Bianji Weiyuanhui				
DT	Journal				
LA	Chinese				
L12	ANSWER 56 OF 70	CAPLUS	COPYRIGHT 2003 ACS on STN		
AN	2001:335842	CAPLUS			
DN	135:251531				
TI	Comparative study of a novel nucleoside analogue (Troxatyl, troxacicabine, BCH-4556) and AraC against leukemic human tumor xenografts expressing high or low cytidine deaminase activity				
AU	Gourdeau, Henriette; Bibeau, Lucie; Ouellet, France; Custea, Dominique; Bernier, Louise; Bowlin, Terry				
CS	BioChem Pharma Inc., Laval, QC, H7V 4A7, Can.				
SO	Cancer Chemotherapy and Pharmacology (2001), 47(3), 236-240 CODEN: CCPHDZ; ISSN: 0344-5704				
PB	Springer-Verlag				
DT	Journal				
LA	English				
RE.CNT	20	THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT			
L12	ANSWER 57 OF 70	CAPLUS	COPYRIGHT 2003 ACS on STN		
AN	2001:327881	CAPLUS			
DN	135:236001				
TI	Cyclopentenyl cytosine increases the phosphorylation and incorporation into DNA of arabinofuranosyl cytosine in a myeloid leukemic cell-line				
AU	Verschuur, A. C.; Van Gennip, A. H.; Leen, R.; Voute, P. A.; Van Kuilenburg, A. B. P.				
CS	Academic Medical Centre, Departments of Pediatrics and Clinical Chemistry, Laboratory of Genetic Metabolic Diseases, University of Amsterdam, Amsterdam, 1100 DE, Neth.				
SO	Advances in Experimental Medicine and Biology (2000), 486(Purine and Pyrimidine Metabolism in Man X), 311-317 CODEN: AEMBAP; ISSN: 0065-2598				
PB	Kluwer Academic/Plenum Publishers				
DT	Journal				

LA English

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
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L12 ANSWER 58 OF 70 CAPLUS COPYRIGHT 2003 ACS on STN
AN 2001:57970 CAPLUS
DN 134:261001
TI Simultaneous treatment with 1-.beta.-D-arabinofuranosylcytosine and daunorubicin induces cross-resistance to both drugs due to a combination-specific mechanism in HL60 cells
AU Takemura, Haruyuki; Urasaki, Yoshimasa; Yoshida, Akira; Fukushima, Toshihiro; Ueda, Takanori
CS First Department of Internal Medicine, Fukui Medical University, Fukui, 910-1193, Japan
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CS Department of Internal Medicine III, University Hospital Grosshadern, Ludwig-Maximilians University, Munich, 81377, Germany
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SO Anderson Cancer Center, Houston, TX, 77030, USA
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